

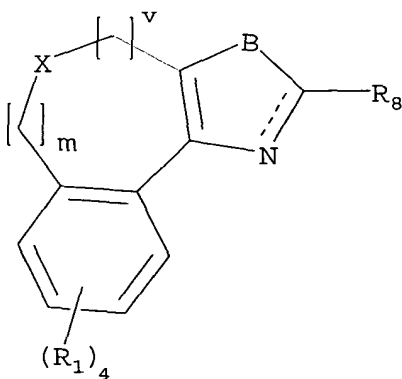
Amendments to the claims:

Please cancel claims 1, 15, 22 and 43 to 46 without disclaimer or prejudice to applicants' right to pursue the subject matter of these claims in a future continuation or divisional application.

Claims 1-46 (canceled)

Please add new claims 47 to 52 as follows:

47. (New) A method of treating a subject suffering from an abnormality wherein the abnormality is alleviated by decreasing the activity of a human Y5 receptor comprising administering to the subject a therapeutically effective amount of a compound having the following structure:



wherein each  $R_1$  is independently H, F, Cl, Br, -CN, -OH, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -SO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>OR<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>CONR<sub>5</sub>R<sub>6</sub>, -(CH<sub>2</sub>)<sub>n</sub>NR<sub>5</sub>COR<sub>5</sub>, perfluoroalkyl, polyfluoroalkyl, aminoalkyl, or straight chained or branched C<sub>1</sub>-C<sub>7</sub> alkyl;

wherein  $R_5$  is independently H; or straight chained or branched  $C_1$ - $C_7$  alkyl;

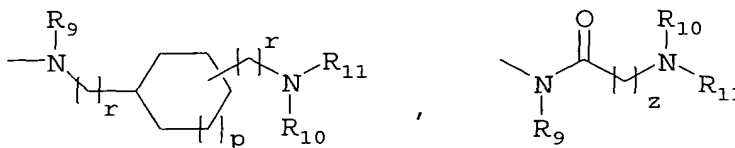
wherein  $R_6$  is independently H; or straight chained or branched  $C_1$ - $C_7$  alkyl;

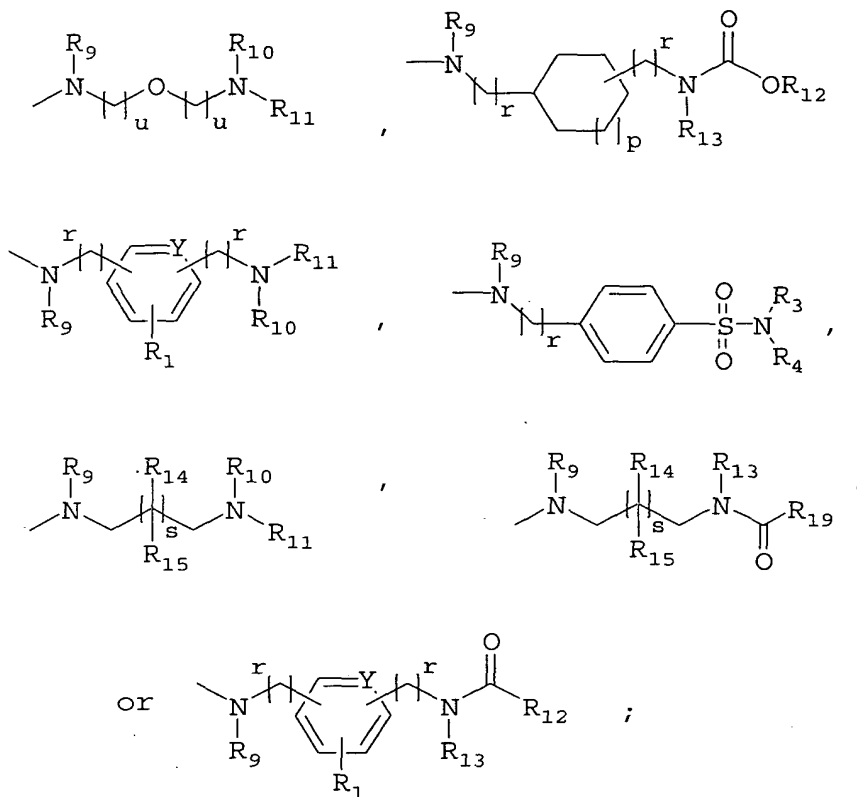
wherein B is O, NH or S;

wherein X is  $CHR_5$ , O or  $NR_5$ ;

wherein each n independently is an integer from 0 to 6 inclusive;

wherein  $R_8$  is





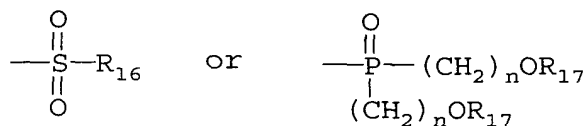
wherein  $Y$  is C or N;

wherein  $R_7$  is independently straight chained or branched  $C_1$ - $C_7$  alkyl;

wherein  $R_9$  is independently H; or straight chained or branched  $C_1$ - $C_4$  alkyl;

wherein  $R_{10}$  is independently H; or straight chained or branched  $C_1$ - $C_4$  alkyl;

wherein  $R_{11}$  is



wherein  $\text{R}_{12}$  is H, straight chained or branched  $\text{C}_1$ - $\text{C}_7$  alkyl,  $(\text{CH}_2)_n\text{OR}_{17}$ , or  $\text{O}(\text{CH}_2)_u\text{OR}_{17}$ ; provided that when X is O,  $\text{R}_{12}$  cannot be methyl;

wherein  $\text{R}_{13}$  is independently H;  $-(\text{CH}_2)_u\text{OR}_5$ ;  $-(\text{CH}_2)_t\text{CONR}_5\text{R}_6$ ;  $-(\text{CH}_2)_u\text{NR}_5\text{COR}_5$ ;  $-(\text{CH}_2)_t\text{COR}_7$ ;  $-(\text{CH}_2)_t\text{CO}_2\text{R}_5$ ;  $-(\text{CH}_2)_u\text{NR}_5\text{R}_6$ ;  $-(\text{CH}_2)_u\text{CN}$ ; straight chained or branched  $\text{C}_1$ - $\text{C}_7$  alkyl;  $\text{C}_1$ - $\text{C}_7$  alkyl in which the  $\text{C}_2$ - $\text{C}_7$  atoms may be optionally substituted with one or more F or Cl;  $\text{C}_3$ - $\text{C}_7$  cycloalkyl- $\text{C}_1$ - $\text{C}_7$  alkyl; straight chained or branched  $\text{C}_2$ - $\text{C}_7$  alkenyl or alkynyl; or  $\text{C}_3$ - $\text{C}_7$  cycloalkyl; phenyl or  $\text{C}_1$ - $\text{C}_6$  phenylalkyl; wherein the phenyl or  $\text{C}_1$ - $\text{C}_6$  phenylalkyl may be substituted with one or more of F, Cl,  $-\text{CN}$ ,  $-\text{NO}_2$ ,  $-\text{NR}_5\text{R}_6$ ,  $-\text{SO}_2\text{R}_5$ ,  $-(\text{CH}_2)_n\text{COR}_7$ ,  $-(\text{CH}_2)_n\text{OR}_5$ ,  $-(\text{CH}_2)_n\text{CONR}_5\text{R}_6$ ,  $-(\text{CH}_2)_n\text{NR}_5\text{COR}_5$ ,  $-(\text{CH}_2)_n\text{CO}_2\text{R}_5$ ,  $-(\text{CH}_2)_n\text{SO}_2\text{NR}_5\text{R}_6$ , straight chained or branched  $\text{C}_1$ - $\text{C}_7$  alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl;

or  $\text{R}_{12}$  and  $\text{R}_{13}$  together with the amide linkage to which they are attached are pyrrolidinonyl, piperidonyl, or oxazolidinonyl; provided that when X is O,  $\text{R}_{12}$  and  $\text{R}_{13}$  cannot be oxazolidinonyl;

wherein  $\text{R}_{14}$  is H; straight chained or branched  $\text{C}_1$ - $\text{C}_4$  alkyl; F; or  $-(\text{CH}_2)_r\text{OR}_5$ ;

wherein  $R_{15}$  is H, straight chained or branched  $C_1$ - $C_4$  alkyl, or F;

with the proviso that when  $R_{14}$  is -OH,  $R_{15}$  cannot be F;

wherein  $R_{16}$  is  $NR_3R_4$ ; perfluoroalkyl, unsubstituted straight chained or branched  $C_1$ - $C_7$  alkyl; substituted straight chained or branched  $C_2$ - $C_7$  alkyl, wherein the  $C_2$ - $C_7$  alkyl may be substituted with one or more of F, Cl, -CN,  $-NR_5R_6$ ,  $-SO_2R_5$ ,  $-(CH_2)_nCOR_7$ ,  $-(CH_2)_nOR_5$ ,  $-(CH_2)_nCONR_5R_6$ ,  $-(CH_2)_nNR_5COR_5$ ,  $-(CH_2)_nCO_2R_5$ ,  $-(CH_2)_nOCF_3$ , perfluoroalkyl, polyfluoroalkyl, or aminoalkyl, straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl, or  $C_3$ - $C_7$  cycloalkyl or cycloalkenyl; phenyl, heteroaryl or  $C_1$ - $C_7$  phenylalkyl, wherein the phenyl, heteroaryl or  $C_1$ - $C_7$  phenylalkyl may be substituted with one or more of F, Cl, Br, -CN,  $-NO_2$ ,  $-NR_5R_6$ ,  $-(CH_2)_nNR_5COR_5$ ,  $-SO_2R_5$ ,  $-(CH_2)_nCOR_7$ ,  $-(CH_2)_nOR_5$ ,  $-(CH_2)_nCONR_5R_6$ ,  $-(CH_2)_nCO_2R_5$ ,  $-(CH_2)_nSO_2NR_5R_6$ , ethylenedioxy, methylenedioxy, straight chained or branched  $C_1$ - $C_7$  alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl, straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl, or  $C_3$ - $C_7$  cycloalkyl or cycloalkenyl; quinolinyl, 1-naphthyl, 2-naphthyl, or 2,1,3-benzothiadiazolyl; wherein the quinolinyl, 1-naphthyl, 2-naphthyl, or 2,1,3-benzothiadiazolyl may be substituted with one or more of F, Cl, Br, -CN,  $-NO_2$ ,  $-NR_5R_6$ ,  $-(CH_2)_nNR_5COR_5$ ,  $-SO_2R_5$ ,  $-(CH_2)_nCOR_7$ ,  $-(CH_2)_nOR_5$ ,  $-(CH_2)_nCONR_5R_6$ ,  $-(CH_2)_nCO_2R_5$ ,  $-(CH_2)_nSO_2NR_5R_6$ , ethylenedioxy, methylenedioxy, straight chained or branched  $C_1$ - $C_7$  alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl;

with the proviso that when X is O and R<sub>8</sub> is NR<sub>9</sub>(CH<sub>2</sub>)<sub>u</sub>O(CH<sub>2</sub>)<sub>u</sub>NR<sub>10</sub>R<sub>11</sub>, R<sub>16</sub> cannot be methyl;

wherein R<sub>3</sub> is independently H; -(CH<sub>2</sub>)<sub>u</sub>OR<sub>5</sub>; -(CH<sub>2</sub>)<sub>t</sub>CONR<sub>5</sub>R<sub>6</sub>; -(CH<sub>2</sub>)<sub>u</sub>NR<sub>5</sub>COR<sub>5</sub>; -(CH<sub>2</sub>)<sub>t</sub>COR<sub>7</sub>; -(CH<sub>2</sub>)<sub>t</sub>CO<sub>2</sub>R<sub>5</sub>; -(CH<sub>2</sub>)<sub>u</sub>NR<sub>5</sub>R<sub>6</sub>; -(CH<sub>2</sub>)<sub>u</sub>CN; straight chained or branched C<sub>1</sub>-C<sub>7</sub> alkyl; straight chained or branched C<sub>2</sub>-C<sub>7</sub> alkenyl or alkynyl; or C<sub>3</sub>-C<sub>7</sub> cycloalkyl or cycloalkenyl; phenyl, C<sub>1</sub>-C<sub>6</sub> phenylalkyl, or C<sub>1</sub>-C<sub>6</sub> heteroarylalkyl; wherein the phenyl, C<sub>1</sub>-C<sub>6</sub> phenylalkyl or C<sub>1</sub>-C<sub>6</sub> heteroarylalkyl may be substituted with one or more of F, Cl, Br, -CN, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -SO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>COR<sub>7</sub>, -(CH<sub>2</sub>)<sub>n</sub>OR<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>CONR<sub>5</sub>R<sub>6</sub>, -(CH<sub>2</sub>)<sub>n</sub>NR<sub>5</sub>COR<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>CO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>SO<sub>2</sub>NR<sub>5</sub>R<sub>6</sub>, straight chained or branched C<sub>1</sub>-C<sub>7</sub> alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl, straight chained or branched C<sub>2</sub>-C<sub>7</sub> alkenyl or alkynyl, or C<sub>3</sub>-C<sub>7</sub> cycloalkyl or cycloalkenyl;

wherein R<sub>4</sub> is independently H; -(CH<sub>2</sub>)<sub>u</sub>OR<sub>5</sub>; -(CH<sub>2</sub>)<sub>t</sub>CONR<sub>5</sub>R<sub>6</sub>; -(CH<sub>2</sub>)<sub>u</sub>NR<sub>5</sub>COR<sub>5</sub>; -(CH<sub>2</sub>)<sub>t</sub>COR<sub>7</sub>; -(CH<sub>2</sub>)<sub>t</sub>CO<sub>2</sub>R<sub>5</sub>; -(CH<sub>2</sub>)<sub>u</sub>NR<sub>5</sub>R<sub>6</sub>; -(CH<sub>2</sub>)<sub>u</sub>CN; straight chained or branched C<sub>1</sub>-C<sub>7</sub> alkyl; straight chained or branched C<sub>2</sub>-C<sub>7</sub> alkenyl or alkynyl; or C<sub>3</sub>-C<sub>7</sub> cycloalkyl or cycloalkenyl; phenyl or C<sub>1</sub>-C<sub>6</sub> phenylalkyl; wherein the phenyl or C<sub>1</sub>-C<sub>6</sub> phenylalkyl may be substituted with one or more of F, Cl, Br, -CN, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -SO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>COR<sub>7</sub>, -(CH<sub>2</sub>)<sub>n</sub>OR<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>CONR<sub>5</sub>R<sub>6</sub>, -(CH<sub>2</sub>)<sub>n</sub>NR<sub>5</sub>COR<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>CO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>SO<sub>2</sub>NR<sub>5</sub>R<sub>6</sub>, straight chained or branched C<sub>1</sub>-C<sub>7</sub> alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl, straight chained or branched C<sub>2</sub>-C<sub>7</sub> alkenyl or alkynyl, or C<sub>3</sub>-C<sub>7</sub> cycloalkyl or cycloalkenyl;

or  $R_3$  and  $R_4$  taken together with the nitrogen atom to which they are attached are 1-azetidiny1, 1-pyrrolidinyl, 1-piperidinyl, or 1H-azepanyl, wherein the 1-azetidiny1, 1-pyrrolidinyl, 1-piperidinyl, or 1H-azepanyl is substituted with one or more of F, -CN,  $-(CH_2)_nNR_5R_6$ ,  $-SO_2R_5$ ,  $-(CH_2)_nCOR_7$ ,  $-(CH_2)_nOR_5$ ,  $-(CH_2)_nCONR_5R_6$ ,  $-(CH_2)_nNR_5COR_5$ ,  $-(CH_2)_nCO_2R_5$ , straight chained or branched  $C_1$ - $C_7$  alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl, straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl, or  $C_3$ - $C_7$  cycloalkyl or cycloalkenyl; or phenyl or heteroaryl; wherein if  $-(CH_2)_nNR_5R_6$ ,  $-(CH_2)_nOR_5$ , or  $-(CH_2)_nNR_5COR_5$  are in the 2-position, then n is not 0; wherein the phenyl or heteroaryl may be substituted with one or more of F, Cl, Br, -CN,  $-NO_2$ ,  $-NR_5R_6$ ,  $-SO_2R_5$ ,  $-(CH_2)_nCOR_7$ ,  $-(CH_2)_nOR_5$ ,  $-(CH_2)_nCONR_5R_6$ ,  $-(CH_2)_nNR_5COR_5$ ,  $-(CH_2)_nCO_2R_5$ ,  $-(CH_2)_nSO_2NR_5R_6$ , straight chained or branched  $C_1$ - $C_7$  alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl, straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl, or  $C_3$ - $C_7$  cycloalkyl or cycloalkenyl;

or  $R_3$  and  $R_4$  taken together with the nitrogen atom to which they are attached are morpholinyl, thiomorpholinyl, [1,4]oxazepanyl, [1,4]thiazepanyl, piperazinyl, or [1,4]diazepanyl is optionally substituted with straight chained or branched  $C_1$ - $C_5$  alkyl or  $(CH_2)_tOR_5$ ; and wherein the nitrogen atom of the piperazinyl or [1,4] diazepanyl ring may be optionally substituted with  $-(CH_2)_uOR_5$ ,  $-COR_5$ ; straight chained or branched  $C_1$ - $C_5$  alkyl; or phenyl; wherein the phenyl may be substituted with one or more of F, Cl, Br, -CN,  $-NO_2$ ,  $-NR_5R_6$ ,  $-(CH_2)_nOR_5$ , straight chained or

branched  $C_1$ - $C_3$  alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl;

wherein  $R_{17}$  is H, straight chained or branched  $C_1$ - $C_4$  alkyl, perfluoroalkyl, or polyfluoroalkyl;

wherein  $R_{19}$  is  $-(CH_2)_nOR_5$ ,  $-NR_5R_6$ , phenyl or heteroaryl, wherein the phenyl or heteroaryl may be substituted with one or more of F, Cl, Br,  $-CN$ ,  $-NO_2$ ,  $-NR_5R_6$ ,  $-(CH_2)_nNR_5COR_5$ ,  $-SO_2R_5$ ,  $-(CH_2)_nCOR_7$ ,  $-(CH_2)_nOR_5$ ,  $-(CH_2)_nCONR_5R_6$ ,  $-(CH_2)_nC_2OR_5$ ,  $-(CH_2)_nSO_2NR_5R_6$ , ethylenedioxy, methylenedioxy, straight chained or branched  $C_1$ - $C_7$  alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl, straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl, or  $C_3$ - $C_7$  cycloalkyl or cycloalkenyl;

wherein m is 0 or 1;

wherein each p independently is an integer from 0 to 2 inclusive;

wherein each r independently is an integer from 0 to 3 inclusive;

wherein each s independently is an integer from 1 to 6 inclusive;

wherein t is an integer from 1 to 4 inclusive;



wherein each u independently is an integer from 2 to 4 inclusive;

wherein v is 1 or 2;

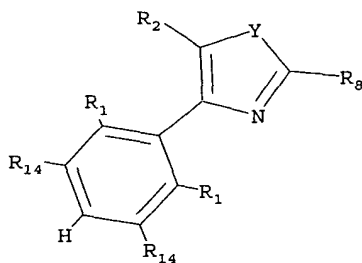
with the proviso that when v is 2, m is 0;

wherein z is an integer from 2 to 7;

or a pharmaceutically acceptable salt thereof to the subject, thereby alleviating the abnormality.

48. (New) The method of claim 47, wherein the abnormality is an eating disorder, obesity, bulimia nervosa, a sexual disorder, a reproductive disorder, depression, an epileptic seizure, hypertension, cerebral hemorrhage, congestive heart failure, or a sleep disturbance.

49. (New) A method of treating a subject suffering from an abnormality wherein the abnormality is alleviated by decreasing the activity of a human Y5 receptor comprising administering to the subject a therapeutically effective amount of a compound having the following structure:



wherein Y is O, S or NH;

wherein each  $R_{14}$  is independently is H, F, Cl, Br, -CN, -OH, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -SO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>OR<sub>5</sub>, -SO<sub>2</sub>C<sub>6</sub>H<sub>5</sub>, -SO<sub>2</sub>NR<sub>5</sub>R<sub>6</sub>, -C<sub>6</sub>H<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>CONR<sub>5</sub>R<sub>6</sub>, -(CH<sub>2</sub>)<sub>n</sub>NR<sub>5</sub>COR<sub>5</sub>, ethylenedioxy, methylenedioxy, perfluoroalkyl, polyfluoroalkyl, aminoalkyl, or straight chained or branched C<sub>1</sub>-C<sub>7</sub> alkyl; or phenyl, heteroaryl or C<sub>1</sub>-C<sub>7</sub> phenylalkyl; wherein the phenyl, heteroaryl or C<sub>1</sub>-C<sub>7</sub> phenylalkyl may be substituted with one or more of F, Cl, Br, -CF<sub>3</sub>, -CN, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -SO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>OR<sub>5</sub>, or straight chained or branched C<sub>1</sub>-C<sub>4</sub> alkyl; provided that if one  $R_{14}$  is phenyl, heteroaryl or C<sub>1</sub>-C<sub>7</sub> phenylalkyl, the other  $R_{14}$  is H;

wherein each  $R_1$  independently is H, F, Cl, Br, -CN, -OH, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -SO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>OR<sub>5</sub>, -SO<sub>2</sub>C<sub>6</sub>H<sub>5</sub>, -SO<sub>2</sub>NR<sub>5</sub>R<sub>6</sub>, -C<sub>6</sub>H<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>CONR<sub>5</sub>R<sub>6</sub>, -(CH<sub>2</sub>)<sub>n</sub>NR<sub>5</sub>COR<sub>5</sub>, ethylenedioxy, methylenedioxy, perfluoroalkyl, polyfluoroalkyl, aminoalkyl, or straight chained or branched C<sub>1</sub>-C<sub>7</sub> alkyl; or phenyl, heteroaryl or C<sub>1</sub>-C<sub>7</sub> phenylalkyl; wherein the phenyl, heteroaryl or C<sub>1</sub>-C<sub>7</sub> phenylalkyl may be substituted with one or more of F, Cl, Br, -CF<sub>3</sub>, -CN, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -SO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>OR<sub>5</sub>, or straight chained or branched C<sub>1</sub>-C<sub>4</sub> alkyl;

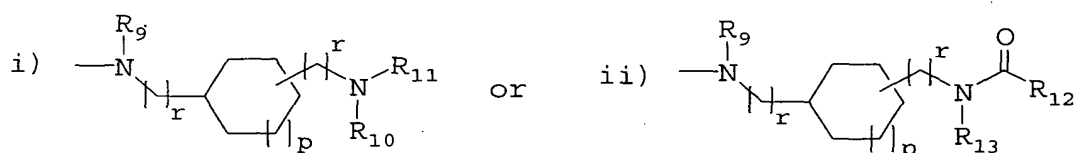
wherein  $R_2$  is H, straight chained or branched C<sub>1</sub>-C<sub>4</sub> alkyl; -(CH<sub>2</sub>)<sub>t</sub>OR<sub>5</sub>, phenyl, optionally substituted with one or more of F, Cl, Br, -CF<sub>3</sub>, -CN, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -SO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>OR<sub>5</sub>, or straight chained or branched C<sub>1</sub>-C<sub>4</sub> alkyl;

wherein  $R_5$  is independently H; or straight chained or branched C<sub>1</sub>-C<sub>7</sub> alkyl;

wherein  $R_6$  is independently H; or straight chained or branched  $C_1$ - $C_7$  alkyl;

wherein each  $n$  independently is an integer from 0 to 6 inclusive;

wherein  $R_8$  is

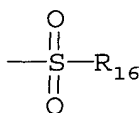


provided that  $R_1$  or  $R_{14}$  cannot be  $-OH$ , when  $R_8$  is (ii);

wherein  $R_9$  is independently H; or straight chained or branched  $C_1$ - $C_4$  alkyl;

wherein  $R_{10}$  is independently H; or straight chained or branched  $C_1$ - $C_4$  alkyl;

wherein  $R_{11}$  is



wherein  $R_{12}$  is H, straight chained or branched  $C_1$ - $C_7$  alkyl; or  $(CH_2)_nOR_{17}$ ;

wherein  $R_{13}$  is independently  $-(CH_2)_uOR_5$ ;  $-(CH_2)_tCONR_5R_6$ ;  $-(CH_2)_uNR_5COR_5$ ;  $-(CH_2)_tCOR_7$ ;  $-(CH_2)_tCO_2R_5$ ;  $-(CH_2)_uNR_5R_6$ ;  $-(CH_2)_uCN$ ; straight chained or branched  $C_1$ - $C_7$  alkyl;  $C_1$ - $C_7$  alkyl in which the  $C_2$ - $C_7$  atoms may be optionally substituted with one or more F or Cl;  $C_3$ - $C_7$  cycloalkyl- $C_1$ - $C_7$  alkyl; straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl; or  $C_3$ - $C_7$  cycloalkyl; phenyl or  $C_1$ - $C_6$  phenylalkyl; wherein the phenyl or  $C_1$ - $C_6$  phenylalkyl may be substituted with one or more of F, Cl,  $-CN$ ,  $-NO_2$ ,  $-NR_5R_6$ ,  $-SO_2R_5$ ,  $-(CH_2)_nCOR_7$ ,  $-(CH_2)_nOR_5$ ,  $-(CH_2)_nCONR_5R_6$ ,  $-(CH_2)_nNR_5COR_5$ ,  $-(CH_2)_nCO_2R_5$ ,  $-(CH_2)_nSO_2NR_5R_6$ , or straight chained or branched  $C_1$ - $C_7$  alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl;

or  $R_{12}$  and  $R_{13}$  together with the amide linkage to which they are attached are pyrrolidinonyl, piperidonyl, or oxazolidinonyl;

wherein  $R_7$  is independently straight chained or branched  $C_1$ - $C_7$  alkyl;

wherein  $R_{16}$  is  $NR_3R_4$ ; perfluoroalkyl, unsubstituted straight chained or branched  $C_1$ - $C_7$  alkyl; substituted straight chained or branched  $C_2$ - $C_7$  alkyl, wherein the  $C_2$ - $C_7$  alkyl may be substituted with one or more of F, Cl,  $-CN$ ,  $-NR_5R_6$ ,  $-SO_2R_5$ ,  $-(CH_2)_nCOR_7$ ,  $-(CH_2)_nOR_5$ ,  $-(CH_2)_nCONR_5R_6$ ,  $-(CH_2)_nNR_5COR_5$ ,  $-(CH_2)_nCO_2R_5$ ,  $-(CH_2)_nOCF_3$ , perfluoroalkyl, polyfluoroalkyl, or aminoalkyl, straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl, or  $C_3$ - $C_7$  cycloalkyl or cycloalkenyl; or phenyl, heteroaryl or  $C_1$ - $C_7$  phenylalkyl; wherein the phenyl, heteroaryl or  $C_1$ - $C_7$  phenylalkyl may be substituted with one

or more of F, Cl, Br, I, -CN, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -(CH<sub>2</sub>)<sub>n</sub>NR<sub>5</sub>COR<sub>5</sub>, -SO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>COR<sub>7</sub>, -(CH<sub>2</sub>)<sub>n</sub>OR<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>CONR<sub>5</sub>R<sub>6</sub>, -(CH<sub>2</sub>)<sub>n</sub>CO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>SO<sub>2</sub>NR<sub>5</sub>R<sub>6</sub>, ethylenedioxy, methylenedioxy, straight chained or branched C<sub>1</sub>-C<sub>7</sub> alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl, straight chained or branched C<sub>2</sub>-C<sub>7</sub> alkenyl or alkynyl, or C<sub>3</sub>-C<sub>7</sub> cycloalkyl or cycloalkenyl; quinolinyl, 1-naphthyl, 2-naphthyl, or 2,1,3-benzothiadiazolyl; wherein the quinolinyl, 1-naphthyl, 2-naphthyl, or 2,1,3-benzothiadiazolyl may be substituted with one or more of F, Cl, Br, I, -CN, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -(CH<sub>2</sub>)<sub>n</sub>NR<sub>5</sub>COR<sub>5</sub>, -SO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>COR<sub>7</sub>, -(CH<sub>2</sub>)<sub>n</sub>OR<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>CONR<sub>5</sub>R<sub>6</sub>, -(CH<sub>2</sub>)<sub>n</sub>CO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>SO<sub>2</sub>NR<sub>5</sub>R<sub>6</sub>, ethylenedioxy, methylenedioxy, or straight chained or branched C<sub>1</sub>-C<sub>7</sub> alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl;

wherein R<sub>3</sub> is independently H; -(CH<sub>2</sub>)<sub>u</sub>OR<sub>5</sub>; -(CH<sub>2</sub>)<sub>t</sub>CONR<sub>5</sub>R<sub>6</sub>; -(CH<sub>2</sub>)<sub>u</sub>NR<sub>5</sub>COR<sub>5</sub>; -(CH<sub>2</sub>)<sub>t</sub>COR<sub>7</sub>; -(CH<sub>2</sub>)<sub>t</sub>CO<sub>2</sub>R<sub>5</sub>; -(CH<sub>2</sub>)<sub>u</sub>NR<sub>5</sub>R<sub>6</sub>; -(CH<sub>2</sub>)<sub>u</sub>CN; straight chained or branched C<sub>1</sub>-C<sub>7</sub> alkyl; straight chained or branched C<sub>2</sub>-C<sub>7</sub> alkenyl or alkynyl; or C<sub>3</sub>-C<sub>7</sub> cycloalkyl or cycloalkenyl; or phenyl, C<sub>1</sub>-C<sub>6</sub> phenylalkyl, or C<sub>1</sub>-C<sub>6</sub> heteroarylalkyl; wherein the phenyl, C<sub>1</sub>-C<sub>6</sub> phenylalkyl, or C<sub>1</sub>-C<sub>6</sub> heteroarylalkyl may be substituted with one or more of F, Cl, Br, -CN, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -SO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>COR<sub>7</sub>, -(CH<sub>2</sub>)<sub>n</sub>OR<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>CONR<sub>5</sub>R<sub>6</sub>, -(CH<sub>2</sub>)<sub>n</sub>NR<sub>5</sub>COR<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>CO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>SO<sub>2</sub>NR<sub>5</sub>R<sub>6</sub>, straight chained or branched C<sub>1</sub>-C<sub>7</sub> alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl, straight chained or branched C<sub>2</sub>-C<sub>7</sub> alkenyl or alkynyl, or C<sub>3</sub>-C<sub>7</sub> cycloalkyl or cycloalkenyl;

wherein  $R_4$  is independently H;  $-(CH_2)_uOR_5$ ;  $-(CH_2)_tCONR_5R_6$ ;  $-(CH_2)_uNR_5COR_5$ ;  $-(CH_2)_tCOR_7$ ;  $-(CH_2)_tCO_2R_5$ ;  $-(CH_2)_uNR_5R_6$ ;  $-(CH_2)_uCN$ ; straight chained or branched  $C_1$ - $C_7$  alkyl; straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl; or  $C_3$ - $C_7$  cycloalkyl or cycloalkenyl; or phenyl or  $C_1$ - $C_6$  phenylalkyl; wherein the phenyl or  $C_1$ - $C_6$  phenylalkyl may be substituted with one or more of F, Cl, Br, -CN, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -SO<sub>2</sub>R<sub>5</sub>,  $-(CH_2)_nCOR_7$ ,  $-(CH_2)_nOR_5$ ,  $-(CH_2)_nCONR_5R_6$ ,  $-(CH_2)_nNR_5COR_5$ ,  $-(CH_2)_nCO_2R_5$ ,  $-(CH_2)_nSO_2NR_5R_6$ , straight chained or branched  $C_1$ - $C_7$  alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl, straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl, or  $C_3$ - $C_7$  cycloalkyl or cycloalkenyl;

or  $R_3$  and  $R_4$  taken together with the nitrogen atom to which they are attached are 1-azetidiny, 1-pyrrolidinyl, 1-piperidinyl, or 1H-azepanyl, wherein the 1-azetidiny, 1-pyrrolidinyl, 1-piperidinyl, or 1H-azepanyl is substituted with one or more of F, -CN,  $-(CH_2)_nNR_5R_6$ , -SO<sub>2</sub>R<sub>5</sub>,  $-(CH_2)_nCOR_7$ ,  $-(CH_2)_nOR_5$ ,  $-(CH_2)_nCONR_5R_6$ ,  $-(CH_2)_nNR_5COR_5$ ,  $-(CH_2)_nCO_2R_5$ , straight chained or branched  $C_1$ - $C_7$  alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl, straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl, or  $C_3$ - $C_7$  cycloalkyl or cycloalkenyl, or phenyl or heteroaryl; wherein if  $-(CH_2)_nNR_5R_6$ ,  $-(CH_2)_nOR_5$ , or  $-(CH_2)_nNR_5COR_5$  are in the 2-position, then n is not 0; wherein the phenyl or heteroaryl may be substituted with one or more of F, Cl, Br, I, -CN, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -SO<sub>2</sub>R<sub>5</sub>,  $-(CH_2)_nCOR_7$ ,  $-(CH_2)_nOR_5$ ,  $-(CH_2)_nCONR_5R_6$ ,  $-(CH_2)_nNR_5COR_5$ ,  $-(CH_2)_nCO_2R_5$ ,  $-(CH_2)_nSO_2NR_5R_6$ , straight chained or branched  $C_1$ - $C_7$  alkyl, perfluoroalkyl, polyfluoroalkyl, or

aminoalkyl, straight chained or branched C<sub>2</sub>-C<sub>7</sub> alkenyl or alkynyl, or C<sub>3</sub>-C<sub>7</sub> cycloalkyl or cycloalkenyl;

or R<sub>3</sub> and R<sub>4</sub> taken together with the nitrogen atom to which they are attached are morpholinyl, thiomorpholinyl, [1,4]oxazepanyl, [1,4]thiazepanyl, piperazinyl, or [1,4]diazepanyl, wherein the morpholinyl, thiomorpholinyl, [1,4]oxazepanyl, [1,4]thiazepanyl, piperazinyl, or [1,4]diazepanyl is optionally substituted with straight chained or branched C<sub>1</sub>-C<sub>5</sub> alkyl or (CH<sub>2</sub>)<sub>t</sub>OR<sub>5</sub>; and wherein the nitrogen atom of the piperazinyl or [1,4]diazepanyl ring may be optionally substituted with -(CH<sub>2</sub>)<sub>u</sub>OR<sub>5</sub>; -COR<sub>5</sub>; straight chained or branched C<sub>1</sub>-C<sub>5</sub> alkyl; or phenyl; wherein the phenyl may be substituted with one or more of F, Cl, Br, -CN, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -(CH<sub>2</sub>)<sub>n</sub>OR<sub>5</sub>, straight chained or branched C<sub>1</sub>-C<sub>3</sub> alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl;

wherein R<sub>17</sub> is straight chained or branched C<sub>1</sub>-C<sub>4</sub> alkyl, perfluoroalkyl, or polyfluoroalkyl;

wherein each p independently is an integer from 0 to 2 inclusive;

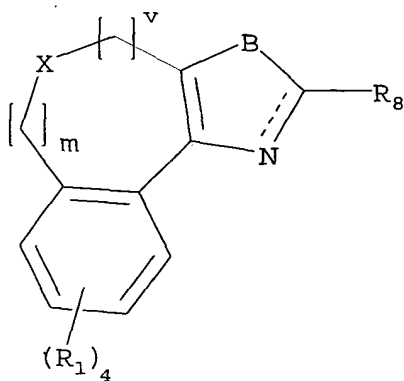
wherein each r independently is an integer from 0 to 3 inclusive;

wherein t is an integer from 1 to 4 inclusive;

wherein each  $u$  independently is an integer from 2 to 4 inclusive;

or a pharmaceutically acceptable salt thereof to the subject, thereby alleviating the abnormality.

50. (New) The method of claim 49, wherein the abnormality is an eating disorder, obesity, bulimia nervosa, a sexual disorder, a reproductive disorder, depression, an epileptic seizure, hypertension, cerebral hemorrhage, congestive heart failure, or a sleep disturbance.
51. (New) A method of treating a subject suffering from an abnormality wherein the abnormality is alleviated by decreasing the activity of a human Y5 receptor comprising administering to the subject a therapeutically effective amount of a compound having the following structure:



wherein each  $R_1$  is independently H, F, Cl, Br, -CN, -OH, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -SO<sub>2</sub>R<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>OR<sub>5</sub>, -(CH<sub>2</sub>)<sub>n</sub>CONR<sub>5</sub>R<sub>6</sub>, -(CH<sub>2</sub>)<sub>n</sub>NR<sub>5</sub>COR<sub>5</sub>, perfluoroalkyl, polyfluoroalkyl, aminoalkyl, or straight chained or branched C<sub>1</sub>-C<sub>7</sub> alkyl;



wherein  $R_5$  is independently H; or straight chained or branched  $C_1$ - $C_7$  alkyl;

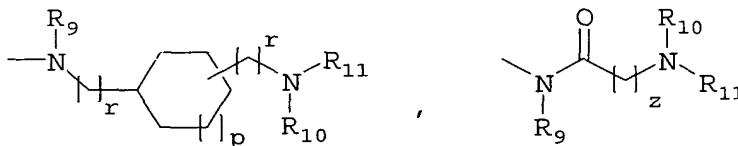
wherein  $R_6$  is independently H; or straight chained or branched  $C_1$ - $C_7$  alkyl;

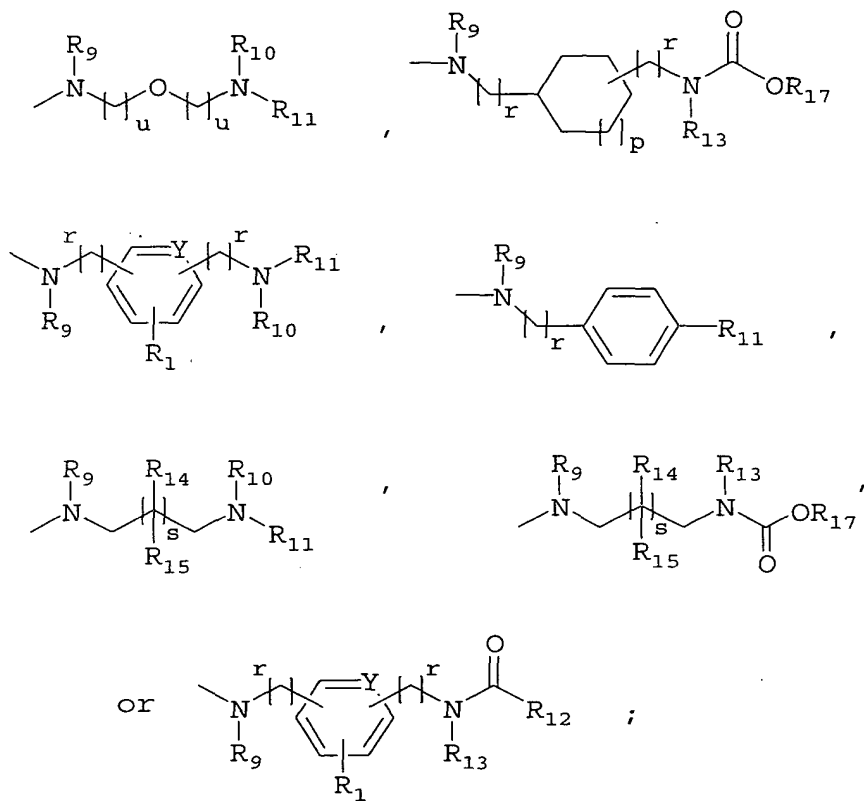
wherein B is O, NH or S;

wherein X is S, SO or  $SO_2$ ;

wherein each n independently is an integer from 0 to 6 inclusive;

wherein  $R_8$  is





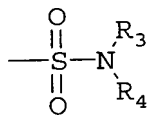
wherein Y is C or N;

wherein  $R_7$  is independently straight chained or branched  $C_1$ - $C_7$  alkyl;

wherein  $R_9$  is independently H; straight chained or branched  $C_1$ - $C_4$  alkyl;

wherein  $R_{10}$  is independently H; or straight chained or branched  $C_1$ - $C_4$  alkyl;

wherein  $R_{11}$  is



wherein  $R_{12}$  is H, straight chained or branched  $C_1$ - $C_7$  alkyl,  $(CH_2)_nOR_{17}$ , or  $O(CH_2)_uOR_{17}$ ;

wherein  $R_{13}$  is independently H;  $-(CH_2)_uOR_5$ ;  $-(CH_2)_tCONR_5R_6$ ;  $-(CH_2)_uNR_5COR_5$ ;  $-(CH_2)_tCOR_7$ ;  $-(CH_2)_tCO_2R_5$ ;  $-(CH_2)_uNR_5R_6$ ;  $-(CH_2)_uCN$ ; straight chained or branched  $C_1$ - $C_7$  alkyl;  $C_1$ - $C_7$  alkyl in which the  $C_2$ - $C_7$  atoms may be optionally substituted with one or more F or Cl;  $C_3$ - $C_7$  cycloalkyl- $C_1$ - $C_7$  alkyl; straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl; or  $C_3$ - $C_7$  cycloalkyl; phenyl or  $C_1$ - $C_6$  phenylalkyl; wherein the phenyl or  $C_1$ - $C_6$  phenylalkyl may be substituted with one or more of F, Cl,  $-CN$ ,  $-NO_2$ ,  $-NR_5R_6$ ,  $-SO_2R_5$ ,  $-(CH_2)_nCOR_7$ ,  $-(CH_2)_nOR_5$ ,  $-(CH_2)_nCONR_5R_6$ ,  $-(CH_2)_nNR_5COR_5$ ,  $-(CH_2)_nCO_2R_5$ ,  $-(CH_2)_nSO_2NR_5R_6$ , straight chained or branched  $C_1$ - $C_7$  alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl;

or  $R_{12}$  and  $R_{13}$  together with the amide linkage to which they are attached are pyrrolidinonyl, piperidonyl, or oxazolidinonyl;

wherein  $R_{14}$  is H; straight chained or branched  $C_1$ - $C_4$  alkyl; F; or  $-(CH_2)_rOR_5$ ;

wherein  $R_{15}$  is H, straight chained or branched  $C_1$ - $C_4$  alkyl, or F;

with the proviso that when  $R_{14}$  is  $-OH$ ,  $R_{15}$  cannot be F;

wherein  $R_3$  is independently H;  $-(CH_2)_uOR_5$ ;  $-(CH_2)_tCONR_5R_6$ ;  $-(CH_2)_uNR_5COR_5$ ;  $-(CH_2)_tCOR_7$ ;  $-(CH_2)_tCO_2R_5$ ;  $-(CH_2)_uNR_5R_6$ ;  $-(CH_2)_uCN$ ; straight chained or branched  $C_1$ - $C_7$  alkyl; straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl; or  $C_3$ - $C_7$  cycloalkyl or cycloalkenyl; phenyl,  $C_1$ - $C_6$  phenylalkyl or  $C_1$ - $C_6$  heteroarylalkyl; wherein the phenyl,  $C_1$ - $C_6$  phenylalkyl, or  $C_1$ - $C_6$  heteroarylalkyl may be substituted with one or more of F, Cl, Br, -CN, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -SO<sub>2</sub>R<sub>5</sub>,  $-(CH_2)_nCOR_7$ ,  $-(CH_2)_nOR_5$ ,  $-(CH_2)_nCONR_5R_6$ ,  $-(CH_2)_nNR_5COR_5$ ,  $-(CH_2)_nCO_2R_5$ ,  $-(CH_2)_nSO_2NR_5R_6$ , straight chained or branched  $C_1$ - $C_7$  alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl, straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl, or  $C_3$ - $C_7$  cycloalkyl or cycloalkenyl;

wherein  $R_4$  is independently H;  $-(CH_2)_uOR_5$ ;  $-(CH_2)_tCONR_5R_6$ ;  $-(CH_2)_uNR_5COR_5$ ;  $-(CH_2)_tCOR_7$ ;  $-(CH_2)_tCO_2R_5$ ;  $-(CH_2)_uNR_5R_6$ ;  $-(CH_2)_uCN$ ; straight chained or branched  $C_1$ - $C_7$  alkyl; straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl; or  $C_3$ - $C_7$  cycloalkyl or cycloalkenyl; phenyl or  $C_1$ - $C_6$  phenylalkyl; wherein the phenyl or  $C_1$ - $C_6$  phenylalkyl may be substituted with one or more of F, Cl, Br, -CN, -NO<sub>2</sub>, -NR<sub>5</sub>R<sub>6</sub>, -SO<sub>2</sub>R<sub>5</sub>,  $-(CH_2)_nCOR_7$ ,  $-(CH_2)_nOR_5$ ,  $-(CH_2)_nCONR_5R_6$ ,  $-(CH_2)_nNR_5COR_5$ ,  $-(CH_2)_nCO_2R_5$ ,  $-(CH_2)_nSO_2NR_5R_6$ , straight chained or branched  $C_1$ - $C_7$  alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl, straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl, or  $C_3$ - $C_7$  cycloalkyl or cycloalkenyl;

or  $R_3$  and  $R_4$  taken together with the nitrogen atom to which they are attached are 1-azetidiny, 1-pyrrolidinyl, 1-

piperidinyl, or 1H-azepanyl, wherein the 1-azetidiny, 1-pyrrolidinyl, 1-piperidinyl, or 1H-azepanyl is substituted with one or more of F, -CN,  $-(CH_2)_nNR_5R_6$ ,  $-SO_2R_5$ ,  $-(CH_2)_nCOR_7$ ,  $-(CH_2)_nOR_5$ ,  $-(CH_2)_nCONR_5R_6$ ,  $-(CH_2)_nNR_5COR_5$ ,  $-(CH_2)_nCO_2R_5$ , straight chained or branched  $C_1$ - $C_7$  alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl, straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl, or  $C_3$ - $C_7$  cycloalkyl or cycloalkenyl, or phenyl or heteroaryl; wherein if  $-(CH_2)_nNR_5R_6$ ,  $-(CH_2)_nOR_5$ , or  $-(CH_2)_nNR_5COR_5$  are in the 2-position, then n is not 0; wherein the phenyl or heteroaryl may be substituted with one or more of F, Cl, Br, -CN,  $-NO_2$ ,  $-NR_5R_6$ ,  $-SO_2R_5$ ,  $-(CH_2)_nCOR_7$ ,  $-(CH_2)_nOR_5$ ,  $-(CH_2)_nCONR_5R_6$ ,  $-(CH_2)_nNR_5COR_5$ ,  $-(CH_2)_nCO_2R_5$ ,  $-(CH_2)_nSO_2NR_5R_6$ , straight chained or branched  $C_1$ - $C_7$  alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl, straight chained or branched  $C_2$ - $C_7$  alkenyl or alkynyl, or  $C_3$ - $C_7$  cycloalkyl or cycloalkenyl;

or  $R_3$  and  $R_4$  taken together with the nitrogen atom to which they are attached are morpholinyl, thiomorpholinyl, [1,4]oxazepanyl, [1,4]thiazepanyl, piperazinyl, or [1,4]diazepanyl, wherein the morpholinyl, thiomorpholinyl, [1,4]oxazepanyl, [1,4]thiazepanyl, piperazinyl, or [1,4]diazepanyl is optionally substituted with  $-(CH_2)_uCOR_5$ ;  $-COR_5$ ;  $-CO_2R_5$ ; straight chained or branched  $C_1$ - $C_5$  alkyl or  $(CH_2)_tOR_5$ ; and wherein the nitrogen atom of the piperazinyl or straight chained or branched  $C_1$ - $C_5$  alkyl; or phenyl; wherein the phenyl may be substituted with one or more of F, Cl, Br, -CN,  $-NO_2$ ,  $-NR_5R_6$ ,  $-(CH_2)_nOR_5$ , straight chained or branched  $C_1$ - $C_3$  alkyl, perfluoroalkyl, polyfluoroalkyl, or aminoalkyl;

wherein  $R_{17}$  is straight chained or branched  $C_1$ - $C_4$  alkyl, perfluoroalkyl, or polyfluoroalkyl;

wherein m is 0 or 1;

wherein each p independently is an integer from 0 to 2 inclusive;

wherein each r independently is an integer from 0 to 3 inclusive;

wherein each s independently is an integer from 1 to 6 inclusive;

wherein t is an integer from 1 to 4 inclusive;

wherein each u independently is an integer from 2 to 4 inclusive;

wherein v is 1 or 2;

with the proviso that when v is 2, m is 0;

wherein z is an integer from 2 to 7;

or a pharmaceutically acceptable salt thereof to the subject, thereby alleviating the abnormality.

52. (New) The method of claim 51, wherein the abnormality is an eating disorder, obesity, bulimia nervosa, a sexual

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disorder, a reproductive disorder, depression, an epileptic seizure, hypertension, cerebral hemorrhage, congestive heart failure, or a sleep disturbance.